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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
09/629,074 07/31/2000		RONALD G CRYSTAL	RONALD G CRYSTAL 205965			
23460	7590 06/05/2002					
LEYDIG VOIT & MAYER, LTD			EXAMI	EXAMINER		
180 NORTH	ENTIAL PLAZA, SUIT STETSON AVENUE	E 4900	BAKER, AN	NE MARIE		
CHICAGO, I	L 60601-6780		ART UNIT	PAPER NUMBER		
			1632	1/		
			DATE MAILED: 06/05/2002	141		

Please find below and/or attached an Office communication concerning this application or proceeding.

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		Application	on No.	Applicant(s)	
•		09/629,07	74	CRYSTAL ET AL.	•
	Office Action Summary	Examin r		Art Unit	
		Anne Bal		1632	
7 Period for f	The MAILING DATE of this communicate Reply	tion app ars on the	cover sheet with the	corr spondence ad	ldr ss
THE MA - Extensio after SIX - If the per - If NO per - Failure to - Any reply	TENED STATUTORY PERIOD FOR ILING DATE OF THIS COMMUNICANS of time may be available under the provisions of 3 (6) MONTHS from the mailing date of this communication for reply specified above is less than thirty (30) date of the specified above, the maximum statutor of the reply is specified above, the maximum statutor reply within the set or extended period for reply will, received by the Office later than three months after that term adjustment. See 37 CFR 1.704(b).	TION. 7 CFR 1.136(a). In no ever cation. ays, a reply within the statu bry period will apply and wi by statute, cause the appl	ent, however, may a reply be story minimum of thirty (30) d Il expire SIX (6) MONTHS fro ication to become ABANDON	timely filed ays will be considered timel m the mailing date of this c IED (35 U.S.C. § 133).	
1)⊠ F	Responsive to communication(s) filed	on <u>08 March 2002</u>	. •		
2a)⊠ T	his action is FINAL . 2b)	☐ This action is	non-final.		
•	ince this application is in condition fo losed in accordance with the practice			_	ne merits is
Disposition	of Claims				
4)⊠ CI	aim(s) <u>1-25</u> is/are pending in the app	olication.			
4a) Of the above claim(s) is/are	withdrawn from co	nsideration.		
5)□ CI	aim(s) is/are allowed.				
6)⊠ CI	aim(s) <u>1-3,6-12,17,23 and 25</u> is/are r	ejected.			
7)⊠ CI	aim(s) <u>4,5,13-16 and 24</u> is/are object	ed to.			
8) CI	aim(s) are subject to restriction	n and/or election re	equirement.		
Application	Papers				
9) 🗌 Th	e specification is objected to by the E	xaminer.			
10) Th	e drawing(s) filed on is/are: a)	accepted or b)	objected to by the Ex	aminer.	
,	Applicant may not request that any object	ion to the drawing(s)	be held in abeyance.	See 37 CFR 1.85(a).	
11) 🔲 Th	e proposed drawing correction filed o	n is: a)∐ a∣	pproved b) disapp	roved by the Examin	ier.
I	f approved, corrected drawings are requir	red in reply to this Of	fice action.		
12) 🗌 Th	e oath or declaration is objected to by	the Examiner.			
Priority und	ler 35 U.S.C. §§ 119 and 120				
13) 🗌 A	cknowledgment is made of a claim for	r foreign priority un	der 35 U.S.C. § 119	(a)-(d) or (f).	
a) <u></u> □	All b) Some * c) None of:				
1.	Certified copies of the priority do	cuments have bee	n received.		
2.	Certified copies of the priority do	cuments have bee	n received in Applica	ation No	
	Copies of the certified copies of to application from the Internation the attached detailed Office action for the action for t	onal Bureau (PCT	Rule 17.2(a)).		Stage
	nowledgment is made of a claim for o		•		al application)
	☐ The translation of the foreign langu				
	knowledgment is made of a claim for	• .	•		
Attachment(s)					
2) Notice o	f References Cited (PTO-892) f Draftsperson's Patent Drawing Review (PTO ion Disclosure Statement(s) (PTO-1449) Pape		' 	ary (PTO-413) Paper No al Patent Application (PT action .	

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DETAILED ACTION

The amendment filed March 8, 2002 (Paper No. 9) has been entered. Claims 1, 6, and 22 have been amended.

Claims 1-25 are pending in the instant application.

The following rejections are reiterated or newly applied and constitute the complete set of rejections being applied to the instant application. Rejections and objections not reiterated from the previous office action are hereby withdrawn.

Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on March 8, 2002 (Paper No. 9) has been entered.

Claim Rejections - 35 USC § 112

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 1-3, 6-12, 17-23, and 25 stand rejected under 35 U.S.C. 112, first paragraph, for reasons of record advanced on pages 4-10 of the Office Action of Paper No. 5 (mailed 4/24/01) and on pages 4-6 of the Office Action of Paper No. 7 (mailed 11/23/01), because the specification, while being enabling for administering either 1) a vector encoding FGF or VEGF operably linked to a promoter or 2) a vector

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encoding FGF or VEGF and a second osteotropic protein each of which is operably linked to a promoter, to a bone or within a tissue immediately surrounding the bone, whereby bone density or formation is enhanced, does not reasonably provide enablement for administration of a first and second nucleic acid to a cell associated with bone, whereby bone density or formation is enhanced, as broadly claimed. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the invention commensurate in scope with these claims.

At page 3, paragraph 3 of the response, Applicants argue that considerable progress has been made on viral targeting and that this technology has been made a part of the instant application by incorporating by reference the text of several patents that concern viral targeting. No specific arguments are offered regarding how these advancements would enable the full scope of the claims, particularly in view of the references cited as evidence that gene therapy is highly unpredictable. However, targeting is not the only issue discussed with regard to lack of an enabling disclosure and, given the state of the art of gene therapy, advances in viral targeting techniques would not be considered sufficient to enable the claimed invention over the full scope because, once delivered, the therapeutic genes must be expressed at a level sufficient to produce the therapeutic effect recited in the claims. The specification and claims contemplate a variety of combinations of angiogenic proteins and osteogenic proteins that can be used in the instant invention. The references cited, including Deonarain (1998), Miller (1995), Verma (1997), and Crystal (1995) highlight the unpredictability in the art of gene therapy and the specific problems associated with obtaining sufficient levels of gene expression to produce a desired effect. Given the unpredictability in the art, the limited guidance in the specification, the limited working examples, undue experimentation would have been required for the skilled artisan to practice the claimed invention over the full scope.

At page 3, paragraph 4 of the response, Applicants argue that hedgehog proteins are osteogenic proteins that can cause bone production *in vivo*. In view of the references cited, the Examiner accepts that

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hedgehog proteins are osteogenic. However, the state scope of enablement for the claimed invention is maintained for the reasons discussed above, and reasons of record.

Thus, the rejection under 35 U.S.C. 112, first paragraph, is maintained for reasons of record.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary.

Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 19-21 stand rejected under 35 U.S.C. 103(a) as being unpatentable over U.S. Patent No. 5,942,496 (Bonadio et al., 8/24/99), for reasons of record advanced on pages 16-17 of the Office Action of Paper No. 5 (mailed 4/24/01) and on page 7 of the Office Action of Paper No. 7 (mailed 11/23/01).

The claims are directed to a viral vector comprising at least one first nucleic acid encoding at least one angiogenic protein and at least one second nucleic acid encoding at least one osteogenic protein.

Bonadio et al. (1999) teaches administration of a nucleic acid encoding FGF and a second osteotropic protein to bone progenitor cells. Claim 33 of Bonadio et al. specifically recites the use of an

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adenovirus vector having a DNA insert comprising one or more osteotropic genes. The instant specification discloses, at page 3, line 11, that FGF is an angiogenic protein. See Claim 40 of Bonadio et al. which clearly shows that an FGF gene can be used in combination with a PTH gene or a BMP gene as recited in Claim 34 from which Claim 40 depends. Both PTH and BMP are disclosed as osteogenic proteins in the instant specification (page 3, lines 18-38).

At page 4, paragraph 2 of the response, Applicants argue that Bonadio et al. does not disclose or suggest a viral vector that has two transgenes, one encoding an angiogenic protein and one encoding an osteogenic protein. However, as discussed above FGF is considered an angiogenic protein and the claims, particularly Claim 40, contemplates using an FGF-encoding nucleic acid in combination with a PTH-encoding gene or a BMP-encoding gene, both osteogenic proteins, as defined in the instant specification.

Thus, the rejection is maintained, for reasons of record.

Conclusion

Claims 4, 5, 13-16, and 24 are objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

No claims are allowable.

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114.

Accordingly, THIS ACTION IS MADE FINAL even though it is a first action after the filing of a request for continued examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b).

Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

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A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Anne-Marie Baker whose telephone number is (703) 306-9155. The examiner can normally be reached Monday through Thursday and alternate Fridays from 10:00 AM to 7:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Deborah Reynolds, can be reached on (703) 305-4051. The fax phone number for the organization where this application or proceeding is assigned is (703) 308-4242.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the patent analyst, Dianiece Jacobs, whose telephone number is (703) 305-3388.

Anne-Marie Baker, Ph.D.

Anne-Marie Baker

PATENT FXAMINER